

IN THE UNITED STATES DISTRICT COURT  
FOR THE DISTRICT OF DELAWARE

	)	
IN RE TRICOR DIRECT PURCHASER	)	
ANTITRUST LITIGATION	)	C.A. No. 05-340 (KAJ)
	)	
	)	(consolidated)
THIS DOCUMENT RELATES TO:	)	
	)	
C.A. No. 05-404 (KAJ)	)	
	)	

**ABBOTT AND FOURNIER'S FIRST AMENDED NOTICE OF VIDEOTAPE  
DEPOSITION OF ECKERD CORPORATION PURSUANT TO RULE 30(B)(6) OF THE  
FEDERAL RULES OF CIVIL PROCEDURE**

To: All Counsel on the Attached Service List

PLEASE TAKE NOTICE that pursuant to Rule 30(b)(6) of the Federal Rules of Civil Procedure, counsel for Defendants shall take the deposition by oral examination of plaintiff Eckerd Corporation ("Eckerd"), on November 8, at the Courtyard Pittsburgh Monroeville, 3962 William Penn Highway, Monroeville, Pennsylvania 15146 (Tel. 412.856.8680), or such other location agreed to by counsel. The deposition will be recorded by videotape as well as stenographically before a Notary Public or other officer authorized to administer oaths, and shall continue from day to day until completed, with such adjournments as to time and place as may be necessary.

NOTICE IS HEREBY GIVEN that pursuant to Fed. R. Civ. P. 30(b)(6), Eckerd is required to designate one or more appropriate persons to testify on its behalf with respect to each of the matters set forth in Exhibit A hereto, and the person(s) so designated shall be required to testify as to each of those matters known or reasonably available to the corporation. You are invited to attend and cross-examine.

FURTHER, Defendants hereby request the production of all documents concerning the topics set forth in Exhibit A that have not already been produced in this litigation.

Production is requested by November 6, 2006.

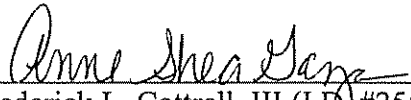
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Dated: November 2, 2006

  
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**DEFINITIONS AND INSTRUCTIONS**

1. The use of any definition for purposes of this Notice shall not be deemed to constitute an agreement or acknowledgement on the part of defendant that such definition is accurate, meaningful or appropriate for any other purpose in this action.

2. “Eckerd” means plaintiff Eckerd Corporation, all parents, subsidiaries, and affiliates thereof, all divisions, predecessors, successors and assigns of each of the foregoing; and all officers, directors, employees, agents, consultants, attorneys and all other persons acting or purporting to act on behalf of, or under the control of, each of the foregoing.

3. “Teva” means counterclaim plaintiffs Teva Pharmaceuticals USA, Inc., Teva Pharmaceutical Industries Ltd., and Novopharm Ltd., all parents, subsidiaries, and affiliates thereof, all divisions, predecessors, successors and assigns of each of the foregoing; and all officers, directors, employees, agents, consultants, attorneys and all other persons acting or purporting to act on behalf of, or under the control of, each of the foregoing.

4. “Impax” means counterclaim plaintiff Impax Laboratories, Inc., all parents, subsidiaries, and affiliates thereof, all divisions, predecessors, successors and assigns of each of the foregoing; and all officers, directors, employees, agents, consultants, attorneys and all other persons acting or purporting to act on behalf of, or under the control of, each of the foregoing.

5. “Fournier” means Fournier Industrie et Santé and Laboratoires Fournier S.A.; all subsidiaries, and affiliates thereof; all divisions, predecessors, successors and assigns of each of the foregoing; and all officers, directors, employees, agents, consultants, attorneys and all

other persons acting or purporting to act on behalf of, or under the control of, each of the foregoing.

6. “Abbott” means Abbott Laboratories; all parents, subsidiaries, and affiliates thereof; all divisions, predecessors, successors and assigns of each of the foregoing; and all officers, directors, employees, agents, consultants, attorneys and all other persons acting or purporting to act on behalf of, or under the control of, each of the foregoing.

7. “McKesson” means McKesson Corporation; all parents, subsidiaries, and affiliates thereof; all divisions, predecessors, successors and assigns of each of the foregoing; and all officers, directors, employees, agents, consultants, attorneys and all other persons acting or purporting to act on behalf of, or under the control of, each of the foregoing.

8. “First Databank” means First Databank, Inc. all parents, subsidiaries, and affiliates thereof; all divisions, predecessors, successors and assigns of each of the foregoing; and all officers, directors, employees, agents, consultants, attorneys and all other persons acting or purporting to act on behalf of, or under the control of, each of the foregoing.

9. “Medi-Span” means Medi-Span of Wolters Kluwer Health; all parents, subsidiaries, and affiliates thereof; all divisions, predecessors, successors and assigns of each of the foregoing; and all officers, directors, employees, agents, consultants, attorneys and all other persons acting or purporting to act on behalf of, or under the control of, each of the foregoing.

10. “Thomson” means Thomson Corporation; all parents, subsidiaries, and affiliates thereof; all divisions, predecessors, successors and assigns of each of the foregoing; and



all officers, directors, employees, agents, consultants, attorneys and all other persons acting or purporting to act on behalf of, or under the control of, each of the foregoing.

11. “TriCor®” means any pharmaceutical product marketed under the trade name “TriCor®” at any time.

12. “Pharmaceutical Product” means any human prescription drug preparation in tablet or capsule form.

13. “Fenofibrate Product” means any human prescription drug preparation in any form, whether or not ultimately offered for sale.

14. “Impax Fenofibrate Product” means any Fenofibrate Product developed, manufactured, marketed, licensed, or sold by Impax.

15. “Teva Fenofibrate Product” means any Fenofibrate Product developed, manufactured, marketed, licensed, or sold by Teva.

16. The term “TriCor®” means any pharmaceutical product marketed under the trade name “TriCor®,” at any time.

17. The term “Lofibra®” means any pharmaceutical product marketed under the trade name “Lofibra®,” at any time.

18. The term “Antara®” means any pharmaceutical product marketed under the trade name “Antara®,” at any time.

19. The term “Triglide®” means any pharmaceutical product marketed under the trade name “Triglide®,” at any time.

20. The term “Lipofen®” means any pharmaceutical product marketed under the trade name “Lipofen®” at any time.

21. The term “document” or “documents” as used herein shall be given the broadest and most inclusive construction provided for under the Federal Rules of Civil Procedure, and includes but is not limited to the following: reports, records, letters, correspondence, telegrams, teletypes, faxes, communications, memoranda, recordings, video, photographs, drawings, books, interoffice communications, bulletins, circulars, pamphlets, brochures, charts, graphs, manuals, minutes, notes, agenda, announcements, instructions, drafts, calendars, diaries, telephone logs, statements, analyses, worksheets, credit memoranda, sales slips, billings or credit statements, ledgers, computer printouts, e-mail, data (including computer diskettes, hard drives, backup disks and tapes), contracts and agreements of any kind. A draft or non-identical copy is a separate document within the meaning of this term.

22. The term “communication” means the transmittal of information by means of documents (as defined above), oral communications, telegrams, voice mail, electronic mail or any other conveyance of information in any form whatsoever, including but not limited to facts, ideas, inquiries or otherwise, regardless of whether published internally or externally.

23. The terms “concern” or “concerning” mean mentioning, identifying, describing, discussing, evidencing, summarizing, explaining, disclosing, recording, showing, containing, setting forth, constituting, comprising, supporting, characterizing, referring to, relating to, and/or regarding, directly or indirectly.

24. As used herein, “and” or “or” shall be considered conjunctively or disjunctively as necessary to make the discovery request inclusive rather than exclusive. The use

of the singular of any word shall include the plural and vice versa, and the use of a verb in any tense or voice shall be construed as the use of that verb in all other tenses and voices, as necessary to bring within the scope of the discovery request all responses that might otherwise be construed as outside its scope.

25. Unless otherwise indicated, the time period applicable to these topics shall be January 1, 1998 to the present.

**EXHIBIT A**

Eckerd is requested to designate one or more officers, directors or managing agents, or other persons who consent to testify on its behalf who have knowledge of the matters set forth herein.

**TOPICS**

1. Eckerd's agreement with McKesson assigning McKesson's claims to Eckerd.
2. Eckerd's records concerning fenofibrate purchases, inventory, and/or returns, including the policies and procedures Eckerd's used to determine the quantities of Fenofibrate Products to purchase, including (1) TriCor®, (2) any Teva Fenofibrate Product, including Lofibra®, (3) Antara®, (4) Triglide®, (5) Lipofen®, and (6) any Impax Fenofibrate Product, including but not limited to the data produced electronically by Eckerd in the file entitled "51998\_1.XLS".
3. The process(es), method(s), strategies, and/or procedures Eckerd proposed, considered, or used for setting or establishing the prices (whether direct or contract, and including rebates, discounts, and/or chargebacks) for any lipid regulating product, including (1) TriCor®, (2) any Teva Fenofibrate Product, including Lofibra®, (3) Antara®, (4) Triglide®, (5) Lipofen®, (6) any Impax Fenofibrate Product, (7) statins, and (8) any other lipid regulating product.
4. Eckerd's ability to control or influence the type or amount of pharmaceutical products demanded by its customers or prescribed for consumers, and the substitutability between any such products, including (1) TriCor®, (2) any Teva Fenofibrate Product, including Lofibra®, (3) Antara®, (4) Triglide®, (5) Lipofen®, (6) any Impax Fenofibrate Product, (7) statins, and (8) any other lipid regulating product, including but not

limited to Eckerd's Generic Fenofibrate Retail Pilot Program, Fenofibrate Pilot Program, Generic Awareness Program, or Compliance and Persistency Program for Eckerd Fenofibrate Patients identified in documents produced by Eckerd Bates labeled Eck 0152-64, Eck 0197-98, Eck 0207-209, Eck 0383-86, Eck 0387-89, and Eck 0445-56.

5. Communications with customers, manufacturers, suppliers, and competitors concerning the availability of fenofibrate products, including (1) TriCor®, (2) any Teva Fenofibrate Product, including Lofibra®, (3) Antara®, (4) Triglide®, (5) Lipofen®, and (6) any Impax Fenofibrate Product, and any analyses of the effect supply shortages had on the price Eckerd's paid for any fenofibrate product.

6. Communications or analyses concerning any comparisons between any one TriCor® formulation on the one hand and (i) any other TriCor® formulation, (ii) any other Fenofibrate Product, (iii) any statin, or (iv) any other lipid regulating product, on the other hand.

7. Eckerd's corporate and personnel organization structure.

8. The methodology used to collect and produce documents in response to Defendants' requests for production of documents and things dated September 30, 2005, June 27, 2006, and August 18, 2006. Such topic includes:

- (i) The document retention orders Eckerd issued to its employees in connection with its obligation to preserve documents relating to this litigation;
- (ii) The selection criteria used to identify the custodians whose files needed to be searched in response to Defendants' requests for production of documents;

- (iii) The identity of all the custodians whose files were searched, including identification of any shared files searched;
- (iv) The criteria used to identify which hard copy documents were responsive to Defendants' requests for production of documents; and
- (v) The criteria used to identify which soft copy documents, including email, were responsive to Defendants' requests for production of documents, including identification of the search terms used to perform electronic searches, and the methodology employed in said electronic searches.

9. The impact, injuries, and/or damages Eckerd alleges it suffered as a result of Defendants' alleged conduct, including the nature and type of any alleged impact, injury, and/or damages (*e.g.*, whether Eckerd is pursuing an overcharge, lost profits, or some other theory of damages).

10. The type and nature of the injunctive relief sought by Eckerd.

**CERTIFICATE OF SERVICE**

I hereby certify that on November 2, 2006, I caused to be served by hand delivery the foregoing document to the following:

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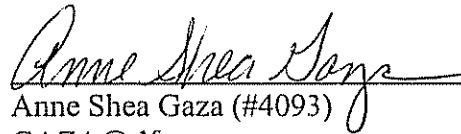
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